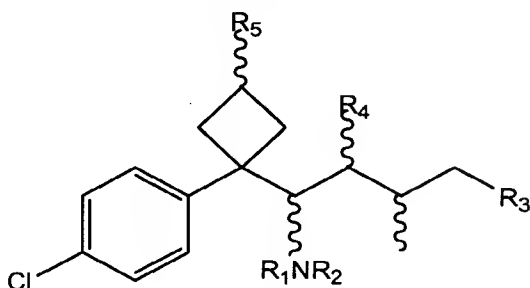


### Amendments to the Claims

The listing of claims will replace all prior versions, and listing of claims in the application.

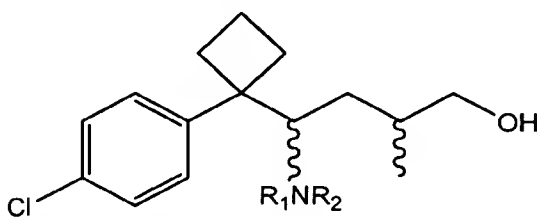
#### Listing of Claims

1. (Original) A compound of the formula:



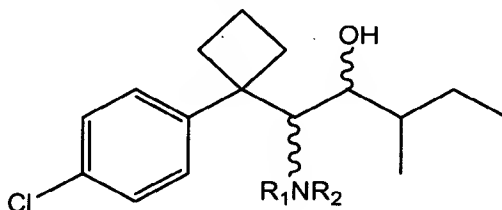
or a pharmaceutically acceptable salt, solvate, hydrate, clathrate, or prodrug thereof, wherein each of R<sub>1</sub> and R<sub>2</sub> is independently lower alkyl or hydrogen, and each of R<sub>3</sub>, R<sub>4</sub>, and R<sub>5</sub> is independently hydrogen, hydroxyl, or alkoxy, provided that: at least one of R<sub>3</sub>, R<sub>4</sub>, and R<sub>5</sub> is not hydrogen; if each of R<sub>1</sub>, R<sub>2</sub>, R<sub>4</sub>, and R<sub>5</sub> is hydrogen and R<sub>3</sub> is hydroxyl, the compound is not racemic; and if each of R<sub>1</sub>, R<sub>2</sub>, R<sub>3</sub>, and R<sub>4</sub> is hydrogen and R<sub>5</sub> is hydroxyl, the compound is not racemic.

2. (Currently Amended) A compound of the formula:



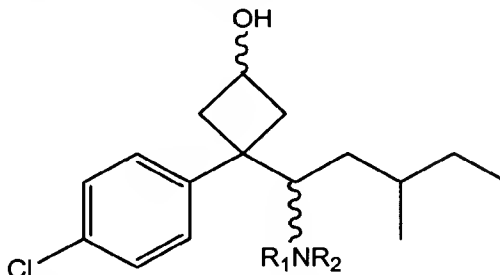
or a pharmaceutically ~~acceptable~~, acceptable salt, solvate, hydrate, clathrate, or prodrug thereof, wherein each of R<sub>1</sub> and R<sub>2</sub> is independently alky or hydrogen, provided that if R<sub>1</sub> and R<sub>2</sub> are both hydrogen, the compound is not racemic.

3. (Currently Amended) A compound of the formula:



or a pharmaceutically ~~acceptable~~, acceptable salt, solvate, hydrate, clathrate, or prodrug thereof, wherein each of R<sub>1</sub> and R<sub>2</sub> is independently alkyl or hydrogen.

4. (Currently Amended) A compound of the formula:



or a pharmaceutically ~~acceptable~~, acceptable salt, solvate, hydrate, clathrate, or prodrug thereof, wherein each of R<sub>1</sub> and R<sub>2</sub> is independently alkyl or hydrogen, provided that if both R<sub>1</sub> and R<sub>2</sub> are hydrogen, the compound is not racemic.

5. (Original) The compound of claim 1, 2, 3, or 4, wherein at least one of R<sub>1</sub> or R<sub>2</sub> is hydrogen.

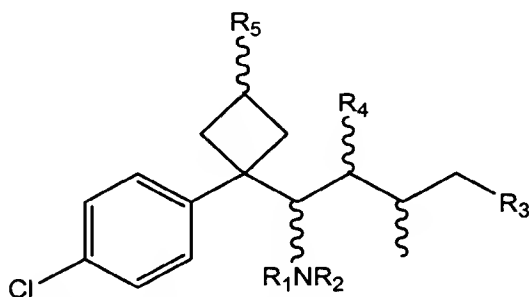
6. (Original) The compound of claim 1, 2, 3, or 4, wherein at least one of R<sub>1</sub> or R<sub>2</sub> is methyl.

7. (Original) The compound of claim 1, 2, 3, or 4, wherein the compound is stereomerically pure.

8. (Original) The compound of claim 1, 2, 3, or 4, wherein the compound is an enantiomeric or diastereomeric mixture that is not a racemic mixture.

Claims 9-31. (Canceled without prejudice)

32. (Original) A pharmaceutical composition comprising a therapeutically or prophylactically effective amount of a racemic or stereomerically pure compound of formula:



or a pharmaceutically acceptable salt, solvate, hydrate, clathrate, or prodrug thereof, wherein each of  $R_1$  and  $R_2$  is independently lower alkyl or hydrogen, and each of  $R_3$ ,  $R_4$ , and  $R_5$  is independently hydrogen, hydroxyl, or alkoxy, provided that at least one of  $R_3$ ,  $R_4$ , and  $R_5$  is not hydrogen.

33. (Original) The pharmaceutical composition of claim 32, wherein if  $R_1$ ,  $R_2$ ,  $R_4$ , and  $R_5$  are each hydrogen and  $R_3$  is hydroxyl, the compound is not racemic, and if  $R_1$ ,  $R_2$ ,  $R_3$ , and  $R_4$  are each hydrogen and  $R_5$  is hydroxyl, the compound is not racemic.

34. (Original) The pharmaceutical composition of claim 32, wherein at least one of  $R_1$  or  $R_2$  is hydrogen and at least one of  $R_3$ ,  $R_4$ , or  $R_5$  is hydroxyl.

35. (Original) The pharmaceutical composition of claim 32, wherein at least one of  $R_1$  or  $R_2$  is methyl and at least one of  $R_3$ ,  $R_4$ , or  $R_5$  is hydroxyl.

36. (Original) The pharmaceutical composition of claim 32, wherein the compound is stereomerically pure.

37. (Original) The pharmaceutical composition of claim 32, wherein the compound is an enantiomeric or diastereomeric mixture that is not racemic.

38. (Original) The pharmaceutical composition of claim 32, wherein the compound is hydroxylated sibutramine or a hydroxylated sibutramine metabolite.

39. (Original) The pharmaceutical composition of claim 38, wherein the compound is hydroxylated in the 1-position.

40. (Original) The pharmaceutical composition of claim 38, wherein the compound is hydroxylated in the 3-position.

41. (Original) The pharmaceutical composition of claim 38, wherein the compound is hydroxylated in the 7-position.

42. (Original) The pharmaceutical composition of claim 32, wherein the pharmaceutical composition is adapted for oral, mucosal, rectal, parenteral, or transdermal administration.

43. (Original) The pharmaceutical composition of claim 32, wherein said composition is lactose-free.

Claims 44-73. (Canceled without prejudice)